

Certificate

acc. to **ISO 13485:2016**

Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes

Certificate Registration No.: 19-1626-Q

TUV USA, Inc. hereby certifies that the quality management system of the company mentioned below is in conformance with **ISO 13485:2016** for Medical Devices-Quality Management Systems-Requirements for Regulatory Purposes.

Wise Plastics Technologies
3810 Stern Avenue
St. Charles, IL 60174, USA

Additional sites covered by QM System: [See Annex I](#)

Scope:

**The Manufacture and Value Added Process Design of Engineered
Injection Molded Products and Assemblies**

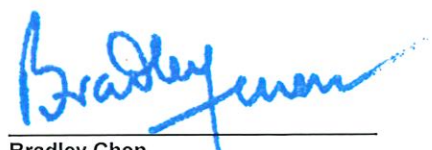
The validity of this certification document can be obtained by contacting the TUV USA, Inc. Office.
TUV USA, Inc. (a Member of the TÜV NORD Group)
215 Main Street, Suite 1, Salem, NH 03079, USA
Tel: 001-603-870-8023, Fax: 001-603-870-8026, Email: medical-usa@tuv-nord.com



Audit Report Reference No.: **19-3705 SA1**
Certificate Initial Issue Date: **2016-05-26**
Current Cycle Start Date: **2019-05-13**
Certificate Revised Date: **2021-05-12**

Effective Date:
2021-05-12 / ed. 3

Valid Until:
2022-05-12



Bradley Chen
Vice President – Medical, Americas
Medical Products Division
TUV USA, Inc.

Annex 1, page 1 of 1
(Annex I MUST be displayed with the main certificate)

Certificate Registration No. : 19-1626-Q / ed. 3
Company Name: Wise Plastics Technologies
Central Office Address: 3810 Stern Avenue, St. Charles, IL 60174, USA



Additional Site(s) covered by the QM System:

Location	Scope of Certification
Headquarters 3810 Stern Avenue St. Charles, IL 60174, USA	The Manufacture and Value Added Process Design of Engineered Injection Molded Products and Assemblies
Site 01 1601 West Hawthorne Ln West Chicago, IL 60185, USA	The Manufacture and Value Added Process Design of Engineered Injection Molded Products and Assemblies

---End of list---

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